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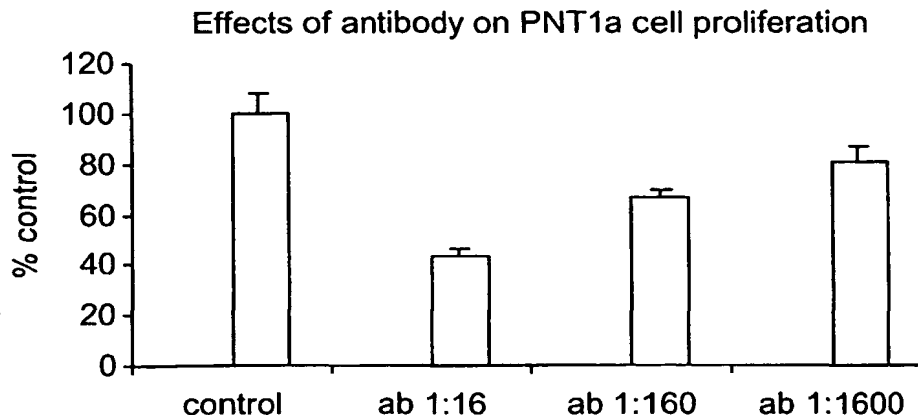
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[Continued on next page]

(54) Title: THERAPEUTIC USES OF MONOCLONAL ANTIBODIES TO THE ANGIOTENSIN-II TYPE-1 RECEPTOR



(57) Abstract: The use of monoclonal antibodies to the angiotensin-II type-I receptor is provided for the treatment of cancer and vascular smooth muscle cell proliferation. Specifically, use is provided of a monoclonal antibody or a fragment thereof to a peptide comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the sequence MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG in the preparation of a medicament for the treatment of cancer or in the preparation of a medicament for the treatment of vascular smooth muscle (VSM) cell proliferation.



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THERAPEUTIC USES OF MONOCLONAL ANTIBODIES TO THE
ANGIOTENSIN-II TYPE-1 RECEPTOR

5 The present invention relates to therapeutic uses of monoclonal antibodies to the angiotensin-II type-I receptor, in particular in the treatment of cancer and vascular smooth muscle cell proliferation.

Angiotensin-II plays a central role in mammalian electrolyte homeostasis and blood pressure control (Peach *Physiol. Rev* **57** 313-370 (1977); Vinson *et al* "The Adrenal Cortex", Prentice Hall, Englefield Heights (1992)). Two main types of angiotensin-II
10 receptors, designated types 1 and 2 (AT1 and AT2), have been recognised, but the majority of the well known actions of angiotensin-II occur via the AT1 subtype (Herblin *et al Am. J. Hypertens.* **4** 299S-302S (1991); Ouali *et al J. Steroid. Biochem. Mol. Biol.* **43** 271-280 (1992)).

15 A monoclonal antibody 6313/G2 to the AT1 receptor subtype (Barker *et al J. Mol. Endocrinol.* **11** 241-245 (1993)) has been used to study the distribution of the receptor (Vinson *et al Mol. Med. Today* **1** 35-38 (1995)). The monoclonal antibody has been suggested for use as a therapeutic agent to control vaso-constriction, for example in
20 the treatment of hypertension or uterine contractions.

The antibody has been used as a specific imaging agent in various tissues, for example laryngeal cancer (Marsigliante *et al Cancer Letters* **110** 19-27 (1996)), kidney (Harrison-Bernard *et al Am. J. Physiol.* **42** F170-F177 (1997); Cheng *et al Am. J. Physiol.* **43** F10-F17 (1998)), and brain (Yang *et al J. Neuroscience* **17** 1660-1669 (1997)). The antibody has been shown to block angiotensin-II induced AT1 receptor
25 internalisation and PKC activation but conversely promotes the calcium response (Kapas *et al Biochem. Biophys. Res. Comm.* **204** 1292-1298 (1994; Vinson *et al J. Endocrinol.* **141** R5-R9 (1994)). The presence of AT1 and At2 receptors in breast
30 tumours has been reported with local production of angiotensin (Inwang *et al Brit. J. Cancer* **75** 1279-1283 (1997); Tahmasebi *et al Eur. J. Cancer* **34** 1777-1782 (1998)).

Monoclonal antibody 6313/G2 is secreted by a hybridoma cell line deposited on 22 July 1993 with the European Collection of Animal Cell Cultures (ECACC), Porton Down, United Kingdom, under the Budapest Treaty, and designated by the accession no. 93072117. The deposit was made by Dr Gavin P Vinson and Dr Stewart Barker,
5 Department of Biochemistry, Queen Mary & Westfield College, Mile End Road, London E1 4NS. The depositor has authorised the applicant to refer to the deposited material in the application and has given his unreserved and irrevocable consent to the deposited material being made available to the public in accordance with Rule 28(1)(d) of the European Patent Convention.

10 The hybridoma cell line produces an antibody that specifically binds to amino acid residues 8 to 17 of the rat vascular smooth muscle AT1 receptor, which sequence is also found in the AT1 receptor of human and bovine cells. The epitope sequence is as follows:

15 EDGIKRIQDD

Or, alternatively expressed as,

20 NH₂-Glu-Asp-Gly-Ile-Lys-Arg-Ile-Gln-Asp-Asp-COOH

It has now been surprisingly found that monoclonal antibodies to the peptide sequence comprising the N-terminal sequence of the angiotensin-II type-1 receptor have additional therapeutic uses in certain medical conditions where such uses were not
25 previously suggested or shown. Furthermore, these therapeutic effects are seen in the ability of the monoclonal antibodies to block the harmful actions of angiotensin-II in the medical conditions concerned whilst preserving the beneficial actions of the molecule. A functionally important role for the entire N-terminal sequence has now been realised.

According to a first aspect of the invention, there is provided the use of a monoclonal antibody or a fragment thereof to a peptide comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the sequence

5 MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG

or a fragment thereof, in the preparation of a medicament for the treatment of cancer.

10 In the above, and throughout this specification, the amino acid residues are designated by the usual IUPAC single letter nomenclature. The single letter designations may be correlated with the classical three letter designations of amino acid residues as follows:

| | | | | |
|----|---------|---------|---------|---------|
| | A = Ala | G = Gly | M = Met | S = Ser |
| | C = Cys | H = His | N = Asn | T = Thr |
| 15 | D = Asp | I = Ile | P = Pro | V = Val |
| | E = Glu | K = Lys | Q = Gln | W = Trp |
| | F = Phe | L = Leu | R = Arg | Y = Tyr |

20 As used herein, the term "peptide" includes oligopeptide or polypeptide and these terms may be used interchangeably.

The peptide will be of at least the minimum size necessary to confer antigenicity: usually it will be of at least six or seven residues, but may be of any suitable length up to, for example, 20 amino acid residues. Preferably, it may be or nine or ten residues. The best
25 peptides may be expected to correspond to topographical surface features of the natural angiotensin-II type-1 receptor molecule, that is to say those features having some three-dimensional feature protruding from or extending into the ambient surface level of the receptor. Preferred peptides correspond to the region 1 to 45, preferably residues 8 to 17.

30

Probably the most simple way of ensuring that at least part of the molecule is antigenically equivalent to the peptide is for that part of the molecule to comprise a

sequence of amino acid residues which is identical to or conformationally similar to the peptide. However, any other way of producing antigenic equivalence may be used: an example is to use an anti-idiotypic antibody or other (even non-proteinaceous) analogue.

5 The invention therefore encompasses the use of monoclonal antibodies against short peptides (preferably of 10 amino acid residues or fewer, but generally of at least 4 or 5 amino acid residues, for example 6 to 8 residues, or 7 to 9 residues) sharing structural homology with the angiotensin-II type-1 receptor.

10 In a preferred embodiment, the invention therefore encompasses the use of a monoclonal antibody against a peptide sequence comprising the amino acid sequence:

EDGIKRIQDD

15 or an active fragment thereof and/or conservative mutant thereof. This sequence is taken from rat angiotensin-II type-1 receptor sequence, residues 8 to 17. Conservative substitutions in this fragment would be D for E, E for D, A for G, L or I, R for K, K for R, N for Q, or any combination of these.

20 The sequence EDGIKRIQDD is fully 100% conserved between the species human, chimpanzee, murine (AT1b) and (AT1b), bovine, canine, ovine, rabbit, and rat (AT1b). Variations are seen in residue 8 for guinea pig which has Q, rat (AT1a) which has D and gerbil which has D. The sequence homologies for these species for the full region 1 to 45 of the N-terminal sequence are shown in Figure 9.

25

Accordingly, a preferred consensus sequence for the peptide corresponding to residues 8 to 17 of the angiotensin-II type -1 receptor has the structure:

EDGIKRIQDD

30

where the following residues may each independently be as follows: residue 8 may be E, D or Q, residue 9 may be D or E, residue 10 may be G or A, residue 11 may be I or A,

residue 12 may be K or R, residue 13 may be R or K, residue 14 may be I or A, residue 15 may be Q or N, and residues 16 and 17 may each either be D or E.

5 The peptide will generally be antigenic and capable of stimulating the production of antibodies which, when administered can be used in the treatment of cancer.

As stated above, an active subfragment of the specified sequence may be used as defined. Active subfragments may consist of or include pentapeptides, including one or more of:

10

TEDGI

EDGIK

DGIKR

GIKRI

15

IKRIQ

KRIQD

RIQDD

IQDDC

20

Active subfragments may also consist of or include hexapeptides, including one or more of:

STEDGI

TEDGIK

25

EDGIKR

DGIKRI

GIKRIQ

IKRIQD

KRIQDD

30

RIQDDC

IQDDCP

Active subfragments may alternatively consist of or include heptapeptides, including one or more of:

5 SSTEDGI
 STEDGIK
 TEDGIKR
 EDGIKRI
 DGIKRIQ
 GIKRIQD
10 IKRIQDD
 KRIQDDC
 RIQDDCP
 IQDDCPK

15 Further, active subfragments may consist of or include octapeptides, including:

 NSSTEDGI
 SSTEDGIK
 STEDGIKR
20 TEDGIKRI
 EDGIKRIQ
 DGIKRIQD
 GIKRIQDD
 IKRIQDDC
25 KRIQDDCP
 RIQDDCPK
 IQDDCPKA

30 Further, active subfragments may consist of or include nonapeptides, including:

 LNSSTEDGI
 NSSTEDGIK

SSTEDGIKR
STEDGIKRI
TEDGIKRIQ
EDGIKRIQD
5 DGIKRIQDD
GIKRIQDDC
IKRIQDDCP
KRIQDDCPK
RIQDDCPKA
10 IQDDCPKAG

Further, active subfragments may consist of or include decapeptides, including:

ILNSSTEDGI
15 LNSSTEDGIK
NSSTEDGIKR
SSTEDGIKRI
STEDGIKRIQ
TEDGIKRIQD
20 EDGIKRIQDD
DGIKRIQDDC
GIKRIQDDCP
IKRIQDDCPK
KRIQDDCPKA
25 RIQDDCPKAG
IQDDCPKAGR

Preferred fragments include those containing some, for example at least four residues of the sequence EDGIKRIQDD.

30

It should be noted that combinations of more than one of the above sequences may be used.

Peptides and other molecules used to prepare monoclonal antibodies for use in accordance with the invention may be rendered antigenic, or presented, in a variety of ways. For preference, an antigenic region (such as a peptide fragment or sub-fragment) in a molecule in accordance with the invention will contain the amino acid sequence of choice linked to a carrier peptide or protein. It is generally preferred to have a plurality, for example 5 to 10, copies of a peptide sequence (for example one or more of the above sequences) linked to the carrier. The carrier can for convenience be a generally large protein, which is inert in material respects, and which is derived from a different species or genus from that associated with the natural growth hormone. Examples of carriers include albumins such as human serum albumin, bovine serum albumin and ovalbumin (although not so many peptides will probably be able to be carried in this last case). Alternatively, keyhole limpet haemocyanin can be used. The carrier will generally preferably come from a different species from that on which the fragment is based.

It is not essential that peptide sequences as described above be linked to albumins: they may be linked to other macromolecules, such as β -galactosidase, especially of bacterial origin.

The invention encompasses the use of monoclonal antibodies to molecules being peptides or having peptide regions which share substantial (e.g. greater than 30%, 50% or even 70%, suitably, 80%, 85%, 90% or 95%) sequence homology with the above peptides. Similarly, conservative amino acid substitutions may not decrease the immunogenicity or antigenicity of peptides. Thus antigenically similar homologues will elicit antibody which binds to angiotensin-II type-1 receptors in the same region as the above peptides define. It is well known that the use of homologues can be a means of circumventing "self" tolerance. Thus the use of the corresponding sequences from other species may be advantageous in this invention.

It is alternatively possible for monoclonal antibodies to be prepared against molecules which are or which comprise peptides to be or to include polymers of sequences as described above. Appropriate sequences can be polymerised either by cross-linking of

two cysteine residues to form disulphide bonds or by using external chemical coupling agents (such as carbodiimide, glutaraldehyde or other dialdehydes or di- (or poly-) functional carboxylic acids. As a further alternative, recombinant DNA techniques could be used to produce a peptide polymer.

5

It should be noted that the chemical coupling (which could for example take place through the agency of lysine residues) and disulphide bond formation are not limited to when the coupling residues are at the end of the sequence: internal residues could also be appropriate. Coupling residues, for example cysteine residues, may be added as desired.

10

It may be found that it is not necessary to couple any of the sequences described above with external peptides. They may be antigenic on their own. In such a case, it may be advisable to select particular adjuvants such as DEAE dextran and Merck 7426.

15

The monoclonal antibodies for used according to the present invention can be prepared by immunising inbred mice by the standard technique of Kohler & Milstein (*Nature* **256** 495-497 (1975)). A peptide corresponding to the epitope sequences described above can be synthesised by any convenient chemical or biological route which is then conjugated to bovine serum albumin (BSA), or another suitable molecule, and then used to immunise the mice. Following a booster injection of the peptide-BSA conjugate, the spleens of the mice are removed, and the spleenocytes combined with mouse myeloma cells. Mixed myeloma-lymphocyte hybrids can then be selected by growth in hypoxanthine, thymidine and aminopterin in an appropriate cell culture medium to inhibit proliferation of non-fused myeloma cells and myeloma hybrids.

25

30

The hybridoma cells can be screened by ELISA for reactivity against the epitope used of the angiotensin-II type-1 receptor by adaptations of the technique described in Engvall *et al Immunochem.* **8** 871 (1991). Alternatively, the antibody capture technique described in Beckmann *et al J. Immunol.* **144** 4212 (1990) may be used. Positive hybridoma cells can be injected intraperitoneally into syngeneic Balb/c mice to produce ascites containing high concentrations of monoclonal antibodies raised

against the epitope used from the angiotensin-II type-1 receptor N-terminal sequence described above. Alternatively, hybridoma cells can be grown *in vitro* in flasks or roller bottles by various techniques. Monoclonal antibodies produced in mouse ascites can be purified by ammonium sulphate precipitation, followed by gel
5 exclusion chromatography. Alternatively, affinity chromatography based upon binding of antibody to protein A or protein G can be used, as can affinity chromatography based upon binding to the epitope used to generate the monoclonal antibody. The monoclonal antibody 6313/G2 was prepared as described in Barker *et al J. Mol. Endocrinol.* **11** 241-245 (1993). Uses of the antibody in the treatment of
10 hypertension and in controlling uterine contractions were described in WO-A-9509186. However, there was no suggestion of any broader utility in other potential therapeutic areas.

The angiotensin-II type-1 receptor of the rat is described in Murphy *et al Nature* **351**
15 233-236 (1992) and the extracellular domain identified as containing at least residues 8 to 17 is represented by the amino acid sequence

EDGIKRIQDD

20 The epitopes from the N-terminal sequence residues 1 to 45, preferably 8 to 17, of the angiotensin-II type-1 receptor described above may be varied modified by amino acid substitution, and/or insertion, and/or deletion such that the overall shape and/or conformation of the epitope is still antigenic.

25 In preferred embodiments of the invention, the monoclonal antibody is 6313/G2. Monoclonal antibody 6313/G2 is secreted by a hybridoma cell line deposited on 22 July 1993 with the European Collection of Animal Cell Cultures (ECACC), Porton Down, United Kingdom, under the Budapest Treaty, and designated by the accession no. 93072117. The hybridoma cell line can suitably be cultured under standard
30 conditions.

In uses of the present invention, the treatment of cancer can include, but is not limited to, inhibition of metastasis, inhibition of binding to matrix proteins of tumour cells, inhibition of invasion by tumour cells and inhibition of tumour cell proliferation. Examples of cancer tumours that may be susceptible to such treatment include, but are not limited to breast cancer and prostate cancer.

In a second aspect of the invention, there is provided the use of a monoclonal antibody to a peptide comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the sequence

10

MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG

15

or a fragment thereof, in the preparation of a medicament for the treatment of vascular smooth muscle (VSM) cell proliferation.

Treatment of vascular smooth muscle cell proliferation may include the treatment of atherosclerosis, a complex disease condition that shows an association with VSM cell proliferation.

The first aspect of the invention therefore also extends to a method for the treatment of cancer comprising administration to a subject in need thereof a therapeutic amount of a monoclonal antibody or a fragment thereof to a peptide comprising the N-terminal portion of the angiotensin-II type-1 receptor or a fragment thereof as defined by the amino acid sequences described above.

25

The second aspect of the invention therefore also extends to a method for the treatment of vascular smooth muscle cell proliferation comprising administration to a subject in need thereof a therapeutic amount of a monoclonal antibody or a fragment thereof to a peptide comprising the N-terminal portion of the angiotensin-II type-1 receptor or a fragment thereof as defined by the amino acid sequence described above.

30

The antibodies used in accordance with the present invention may be formulated for intravenous injection using appropriate pharmaceutically acceptable adjuvants and/or diluents. Injection may be intravenous, intramuscular, intraperitoneal, including subcutaneous injection. Other modes of administration are not excluded, such as for
5 example orally via liposomes, enteric coated capsules and the like.

Suitably, the antibodies used in accordance with the present invention may be humanised antibodies as described in US 4,816,567 and WO 94/10332; or microbodies as described in WO 94/09817; or transgenic antibodies as described in
10 GB-A-2272440. Such synthetic constructs include chimaeric molecules. Thus, for example, uses of humanised (or primatised) antibodies or derivatives thereof are within the scope of the present invention. An example of a humanised antibody is an antibody having human framework regions, but rodent hypervariable regions.

15 In addition to whole antibodies, the present invention includes uses of derivatives of the monoclonal antibodies defined above which are capable of binding to the epitope selected from the N-terminal region of the angiotensin-II type-1 receptor described above. Thus the present invention also includes uses of antibody fragments. Examples of antibody fragments are given by Dougall *et al Tibtech* 12 372-379 (1994).

20 Antibody fragments include, for example, Fab, F(ab')₂ and Fv fragments (Roitt *et al "Immunology"*, Second edition (1989), Churchill Livingstone, London). Fv fragments can be modified to produce a synthetic construct known as a single chain Fv (scFv) molecule. This includes a peptide linker covalently joining V_h and V_l regions which
25 contribute to the stability of the molecule.

Other synthetic constructs include CDR peptides. These are synthetic peptides comprising antigen binding determinants. Peptide mimetics may also be used. These molecules are usually conformationally restricted organic rings which mimic the
30 structure of a CDR loop and which include antigen-interactive side chains. Uses of such molecules able to bind to the desired epitope are therefore also within the scope of the present invention.

In a third aspect of the invention, there is provided the use of a peptide sequence comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the sequence

5

MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG

or a fragment thereof, in the preparation of a medicament for the treatment of cancer.

10 In a fourth aspect of the invention there is provided a vaccine composition comprising a peptide sequence comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the sequence

MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG

15

or a fragment thereof. The vaccine composition may comprise a polypeptide of the above sequence or an antigenic fragment as defined above, optionally conjugated to a carrier protein. Means for rendering such proteins or peptides antigenic are defined above in relation to the earlier aspects of the invention. For example, an albumin protein, such as human serum albumin, bovine serum albumin and ovalbumin. Alternatively, keyhole limpet protein (also sometimes referred to as keyhole limpet haemocyanin) can be used. The carrier will generally be different come from a different species from that on which the fragment is based. Other adjuvants may also be present in the vaccine composition, for example a saponin adjuvant, e.g. a *Quillaja* saponin or a derivative thereof.

20
25

In a particularly preferred embodiment there is provided a method for the inhibition of cancer cell growth, adhesion or invasion comprising:

30 (1) formulating a monoclonal antibody or a fragment thereof to a peptide comprising the N-terminal portion of the angiotensin-II type-1 receptor as defined above, optionally conjugated to a carrier peptide or protein in an appropriate

pharmaceutically acceptable adjuvant and/or diluent, such as for intravenous injection

5 (2) optionally further formulating the monoclonal antibody preparation of (1) as a liposomal or enteric coated capsule formulation

(3) administration of the formulation of (2) or (3) to a population of cancer cells *in vitro* or a subject suffering from cancer.

10 Alternatively, this embodiment may also comprise step (1) and (2) only.

Such embodiments extend to the use of such formulations in the preparation of medicaments for the treatment of cancer, for example, prostate cancer, breast cancer (including breast cancer cell adhesion or invasion).

15

In another preferred embodiment there is provided a method for the inhibition of vascular smooth muscle cell proliferation in which steps (1) to (3) described above are repeated *mutatis mutandis*.

20 Preferred features for the second and subsequent aspects of the invention are as for the first aspect *mutatis mutandis*.

25 The invention will now be further described by way of reference to the following Examples and Figures which are provided for the purposes of illustration only and are not to be construed as being limiting on the invention. Reference is made to a number of Figures in which:

FIGURE 1 shows the effects of antibody on PNT1a cell proliferation.

30 FIGURE 2 shows the effects of antibody on aortic smooth muscle cell proliferation by assaying uptake of tritiated thymidine.

FIGURE 3 shows the results of cell adhesion assay of MCF-7 cells on extracellular matrix protein.

FIGURE 4 shows the results of chemoinvasion assay of MCF-7 cells on extracellular matrix protein.

FIGURE 5 shows results of Western blots in assay of expression of integrins alpha-3 and beta-1 in breast cancer cells.

FIGURE 6 shows antibody 6313/G2 stimulation of calcium responses in MCF-7 cells.

FIGURE 7 shows antibody 6313/G2 stimulation of calcium responses in RASMC.

FIGURE 8 shows a schematic diagram of the actions of angiotensin-II, the site of the monoclonal antibody activation and the site of monoclonal antibody block.

FIGURE 9 shows sequence homologies for the N-terminal sequences of angiotensin-II type-1 receptor from different species, where "X" denotes a missing residue, and "-" denotes an identical residue.

Example 1: Antibody 6313/G2 inhibits cell proliferation in prostate PNT1A cells

The tetrazolium salt, 3-(4,5 dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) is widely used as an indicator for cellular oxidative metabolic activity. On reduction MTT forms an intensely coloured formazan product, which can be measured colorimetrically and thus is often used for the quantitative assessment of cellular viability and proliferation.

PNT1a, a prostate epithelial cell line were seeded into 96 well culture plate at a

concentration of 1000 cells per well. Cells were grown in the presence of RPMI 1640 medium supplemented with 2mM L-glutamine, 1% non-essential amino acids, 2% penicillin and streptomycin, and 1mM sodium pyruvate, 10% fetal calf serum, for two days and subsequently rendered quiescent by incubation in RPMI 1640 serum free medium (200µl/well) for 24 hr. Stimulants were then added at appropriate concentration and incubated for 24 hr and 96 hr. Four hours prior to end of incubation, 20µl of filtered (0.2µm pore size) 5mg/ml solution (in RPMI media) MTT was added to each well and the incubation continued at 37°C. At the appropriate time point, 200µl of DMSO, followed by 25 µl of Sorensen's glycine buffer (0.1M glycine, 0.1M NaCl adjusted to pH 10.5 with 1M NaOH) was added to each well, mixed thoroughly. After 5 minutes, the absorbance was read at 545nm.

Results are shown in Figure 1 in which purified antibody (ab) was added to PNT1a cells in culture. Concentrations of antibody were (1:1600) 100nmol/l, (1:160) 1µmol/l, (1:16) 10µmol/l. Inhibition of proliferation was significant ($P < 0.05$ or better) at all concentrations of antibody used.

Example 2: Antibody 6313/G2 inhibits cell proliferation in vascular smooth muscle cells

ASMCs were isolated from rat thoracic and abdominal artery (RASMC) and bovine aorta (BASMC) by the media explant method and cultured over several passages. Segments of both abdominal and thoracic aortas were obtained from rats by careful dissection from killed rats. Segments of aorta were obtained from calves under anaesthesia. The segments of aorta were placed in a depression slide containing tissue culture medium, after which the adventitia and the outer portion of each segment was carefully removed under a dissecting microscope. The remaining inner portion of the tissue and the intima were removed to a separate dissecting dish and washed several times with fresh culture medium. At this point each segment was cut into approximately 1 mm squares and placed on 25 cm² tissue culture flask. The flasks were loosely capped and placed in a humidified CO₂ incubator. After two hours, 4 ml of RPMI-1640 culture medium supplemented with 100 units/ml of penicillin, 100

- 5 $\mu\text{g/ml}$ streptomycin, 4 $\mu\text{mol/L}$ L-glutamine and 20% FBS was carefully added to the flasks without dislodging the tissue. Samples were fed with fresh medium after one week. The cells from the explants were relatively confluent within a period of approximately 2 weeks. They were then rinsed with PBS, and subsequently trypsinized with a solution of 0.125% trypsin and 0.02% EDTA in PBS for 1-2 minutes at 37°C. The resulting suspension of cells was pipetted into 75 cm^2 tissue culture flasks containing 10 ml culture medium and incubated as above. Experiments were performed with cells from passages 3-5.
- 10 A suspension of RASMC (10^5 cells/ml) were prepared on the first day of the experiment using RPMI-1640 supplemented with 20% FBS. One ml of this cell suspension was distributed to each well of a 24-well multiwell dish. The medium was replaced 24 h after the subculture with RPMI-1640 medium. The quiescent (serum-derived) or serum-replete cells were incubated with the appropriate experimental media for 48 hours 4 wells per group. ^3H -methylthymidine (0.1 mCi/ml) 10 μl was added to each well (1 ml medium/well). 24 hours after the addition of radioactive thymidine, media were aspirated and the cultured cells were rinsed 3 times with cold PBS. Cells were then dissolved in 0.5ml of 0.1 N NaOH and a 0.3 ml aliquot was mixed with 3.5 ml of scintillation fluid and, after standing overnight at room temperature, tritium content was assayed in a liquid scintillation counter.
- 20

Results are shown in Figure 2 in which proliferation was stimulated by 10nmol/l angiotensin-II and inhibited by 6313/G2 (10 $\mu\text{mol/L}$, **P<0.01).

25 **Example 3: Antibody 6313/G2 inhibits cell proliferation in breast cancer cells**

- 30 MCF-7 cells (obtained from American Type Culture Collection (ATCC) Manassas, VA 20108, USA) were plated out in 24 well dishes at a density of 5000 cells per well and grown for 24 hours in Eagles's Minimum Essential Medium (MEM) containing 5% fetal bovine serum (FBS). Cells were then incubated for a further 24 hours in serum free medium. Following this, cells were grown in either serum free medium

alone (control wells) or serum free medium with addition to experimental wells of angiotensin II alone (1-10nM), or with antibody 6313/G2. Each treatment was performed in quadruplicate. Cells were then cultured for 24 hours. After 20 hours tritiated thymidine was added to each well (Amersham Pharmacia Biotech, 5 Amersham, UK, 50µCi/ml, sp. activity 5Ci/mmol) and cells were cultured for a further 4 hours. At the end of this period the medium was aspirated and the cultured cells were rinsed three times with ice-cold buffer solution (50mM Tris-HCl, pH 7.4). Cells were then dissolved in 1ml 0.1N NaOH and 0.5ml of this solution was mixed with 3.5ml of scintillation cocktail (toluene scintillator, Packard Bioscience B.V. 10 Groningen, Netherlands) and tritium content was assayed.

Example 4: Antibody 6313 inhibits breast cancer cell adhesion

A cell adhesion assay of MCF-7 cells on extracellular matrix protein was carried out to investigate. Cell culture (flat bottomed) 96 well plates were coated with graded 15 amounts of purified human matrix protein, Collagen type IV (50µg/well). They were left overnight in a laminar flow cabinet to evaporate, at room temperature.

MCF-7 cells (obtained from American Type Culture Collection (ATCC) Manassas, VA 20108, USA) were treated with 6163/G2 antibody for 48 hours. Controls were 20 untreated. Prior to use, each well was treated with BSA (200µg/ml) to eliminate non-specific binding.

500 MCF-7 cells in cell culture medium (DMEM) were added to each well and incubated at 37 °C in a 5% CO₂ environment for 1 hour. Wells were then washed 3 25 times with serum-free DMEM and stained with Diff-quick fix (7 seconds), Diff quick I (7 seconds) and Diff quick II (10 seconds) and washed once with water.

Wells were then viewed under a microscope, and the numbers of adhering cells counted. Antibody 6313/G2 significantly reduced cell adhesion (P<0.05). Results are 30 shown in Figure 3 in which the number of MCF-7 cells adhered to collagen (50µg/well) against the treatments with and without antibody 6313.

Example 5: Antibody 6313 inhibits breast cancer cell invasion

A chemoinvasion assay of MCF-7 cells on extracellular matrix protein was carried out to investigate. 8µm filter inserts were coated with purified human collagen type IV matrix protein and left overnight in a laminar flow cabinet to dry at room temperature.

5 MCF-7 cells were treated with 6163/G2 antibody (hybridoma supernatant) for 48 hours. Control cells were untreated.

Prior to use, BSA (100µg/ml) was added to each well for 1 hour. DMEM preconditioned by incubation with 3T3 fibroblast cells was used as the

10 chemoattractant. The coated inserts were placed in each well to form an upper and a lower chamber. 10,000 MCF-7 cells were added into the upper chamber with the addition of serum free DMEM. Conditioned 3T3 cell medium was placed in the lower compartment. Plates were covered and incubated at 37 °C in a humidified 5% CO₂ environment for 24 hours.

15 After incubation, the cells remaining on the upper surface of the filter were completely removed and the cells that had traversed the collagen and attached to the lower surface of the filter were stained with Diff-Quik and counted. Results are shown in Figure 4 in which the number of MCF-7 cells invaded through collagen

20 (50µu/well) against treatments with and without antibody 6313. Antibody 6313 significantly inhibited cell invasion ($P>0.01$).

Example 6: Effect of antibody 6313 on integrin expression in breast cancer cells

The effect of antibody 6313/G2 on integrin expression was investigated. The results

25 show that antibody 6313 significantly reduces integrins alpha 3 and beta 1 expression in breast cancer cells.

MCF-7 cell lines was treated for 48 hours with antibody 6313/G2, controls were untreated. Cell membrane fractions were prepared and fractionated non-reduced 8%

30 SDS-PAGE gel.

Proteins were then transferred to a nitrocellulose membrane overnight, 30V at 4°C.

Primary and secondary antibodies for the integrins $\alpha 3$ and $\beta 1$ were used to detect these components on the nitrocellulose membranes using established methods for Western blotting. Luminescent bands were developed by incubating the membrane in enhanced chemiluminescence (ECL) western blotting detection reagents for 1 minute
5 by hyper film ECL exposure. Results are shown in Figure 5 in which , C = control, A = antibody tested, other lanes (M) are molecular weight markers.

Example 7: Effect of antibody 6313 on calcium responses in MCF-7 cells and in RASMC

10 The effect of antibody 6313 on calcium responses in MCF-7 cells and in RASMC was investigated. Antibody 6313 was found to stimulate the calcium response in both.

For calcium ion ($[Ca^{2+}]$) measurement, the cells were loaded with $1\mu M$ fura-2 for 30 minutes in medium-modified Krebs-Ringer bicarbonate solution ($3.6mM K^+$, $1.2mM Ca^{2+}$, $0.5mM Mg^{2+}$, $5mM$ Hepes and $20mM HCO^-$) at $37^\circ C$. For simultaneous
15 measurements of measuring the fluorescence of fura-2, the cells plated on coverslips were mounted on the stage of an inverted microscope (Zeiss) in a modified Krebs-Ringer bicarbonate solution. The excitation wavelengths were 340nm and 380nm, and emission was detected at 510nm. Calcium ion concentration ($[Ca^{2+}]$) was
20 calculated from the ration of fluorescence intensities at excitation wavelengths of 340nm and 380nm.

Results are shown in Figure 6 in MCF-7 cells and in RASMC. The vertical arrow indicates the point of application if antibody 6313/G2. The increased ration of
25 fluorescence intensities is proportional to the intracellular calcium ion concentration.

CLAIMS

1. The use of a monoclonal antibody or a fragment thereof to a peptide comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the sequence

5

MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG

or a fragment thereof, in the preparation of a medicament for the treatment of cancer.

10

2. The use of a monoclonal antibody or a fragment thereof to a peptide comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the sequence

MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG

15

or a fragment thereof, in the preparation of a medicament for the treatment of vascular smooth muscle (VSM) cell proliferation.

20

3. A use as claimed in claim 1 or claim 2, in which the monoclonal antibody is raised against the peptide EDGIKRIQDD.

4. A use as claimed in any one of claims 1 to 3, in which the monoclonal antibody is a humanised antibody.

25

5. A use as claimed in any one of claims 1 to 3, in which the monoclonal antibody is 6313/G2 produced by the hybridoma cell line designated by accession no. 93072117.

30

6. A method for the treatment of cancer comprising administration to a subject in need thereof a therapeutic amount of a monoclonal antibody or a fragment thereof to a peptide comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the amino acid sequence

MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG

or a fragment thereof.

5

7. A method for the treatment of vascular smooth muscle cell proliferation comprising administration to a subject in need thereof a therapeutic amount of a monoclonal antibody or a fragment thereof to a peptide comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the amino acid sequence

10

MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG

or a fragment thereof.

15

8. A method as claimed in claim 6 or claim 7, in which the monoclonal antibody is raised against the peptide EDGIKRIQDD

9. A method as claimed in any one of claims 6 to 8, in which the monoclonal antibody is a humanised monoclonal antibody.

20

10. A method as claimed in any one of claims 6 to 8, in which the monoclonal antibody is 6313/G2 produced by the hybridoma cell line designated by accession no. 93072117.

25

11. The use of a peptide sequence comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the sequence

MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG

30

or a fragment thereof, in the preparation of a medicament for the treatment of cancer.

12. A vaccine composition comprising a peptide sequence comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the sequence

MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG

5

or a fragment thereof.

13. A vaccine composition as claimed in claim 11, in which the peptide is conjugated to a carrier protein.

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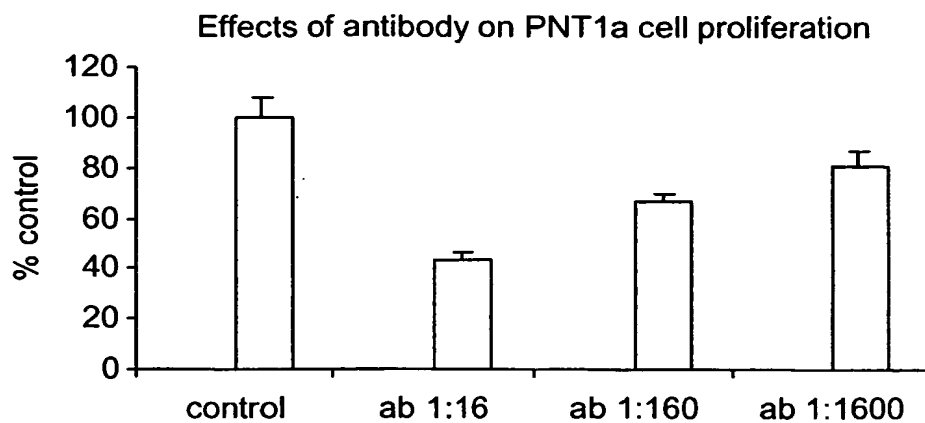


FIG. 1

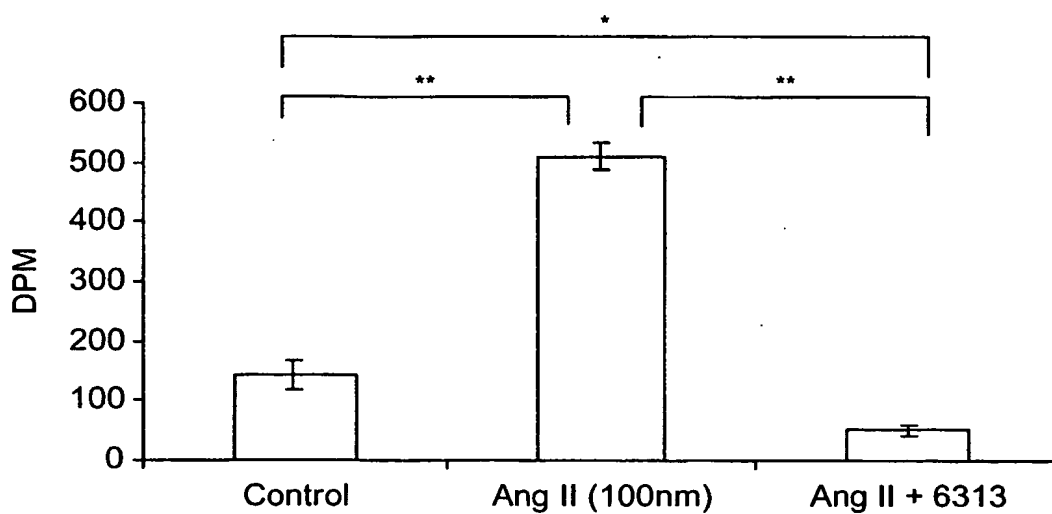


FIG. 2

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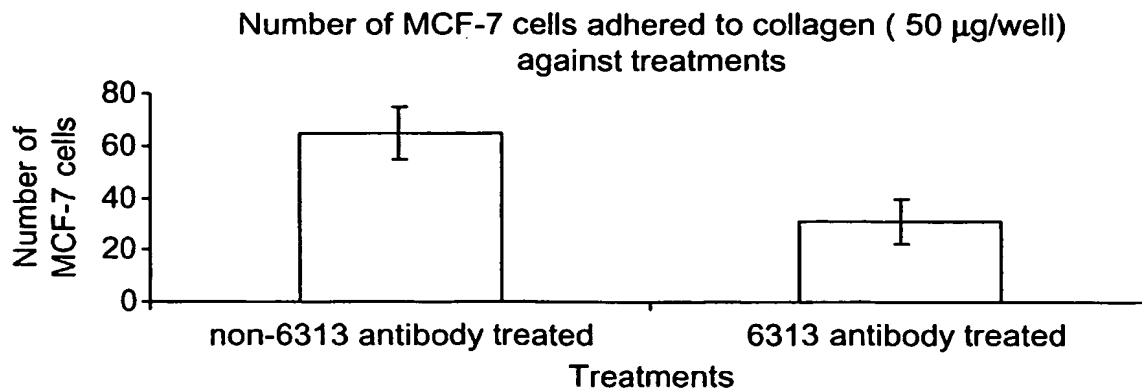


FIG. 3

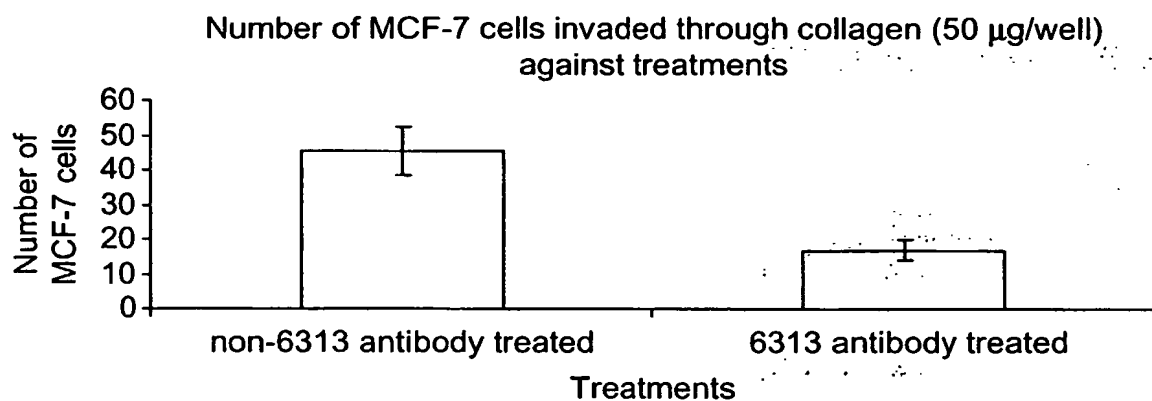


FIG. 4

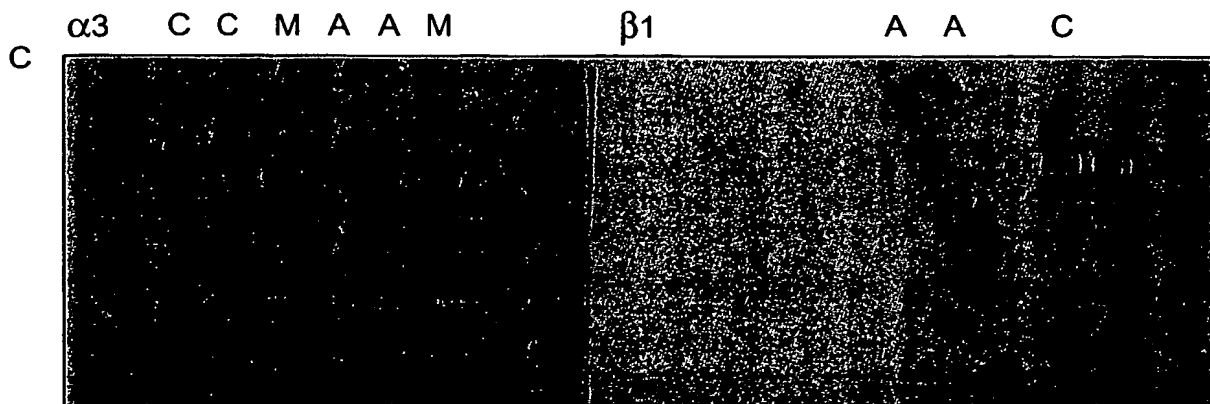


FIG. 5 BEST AVAILABLE COPY

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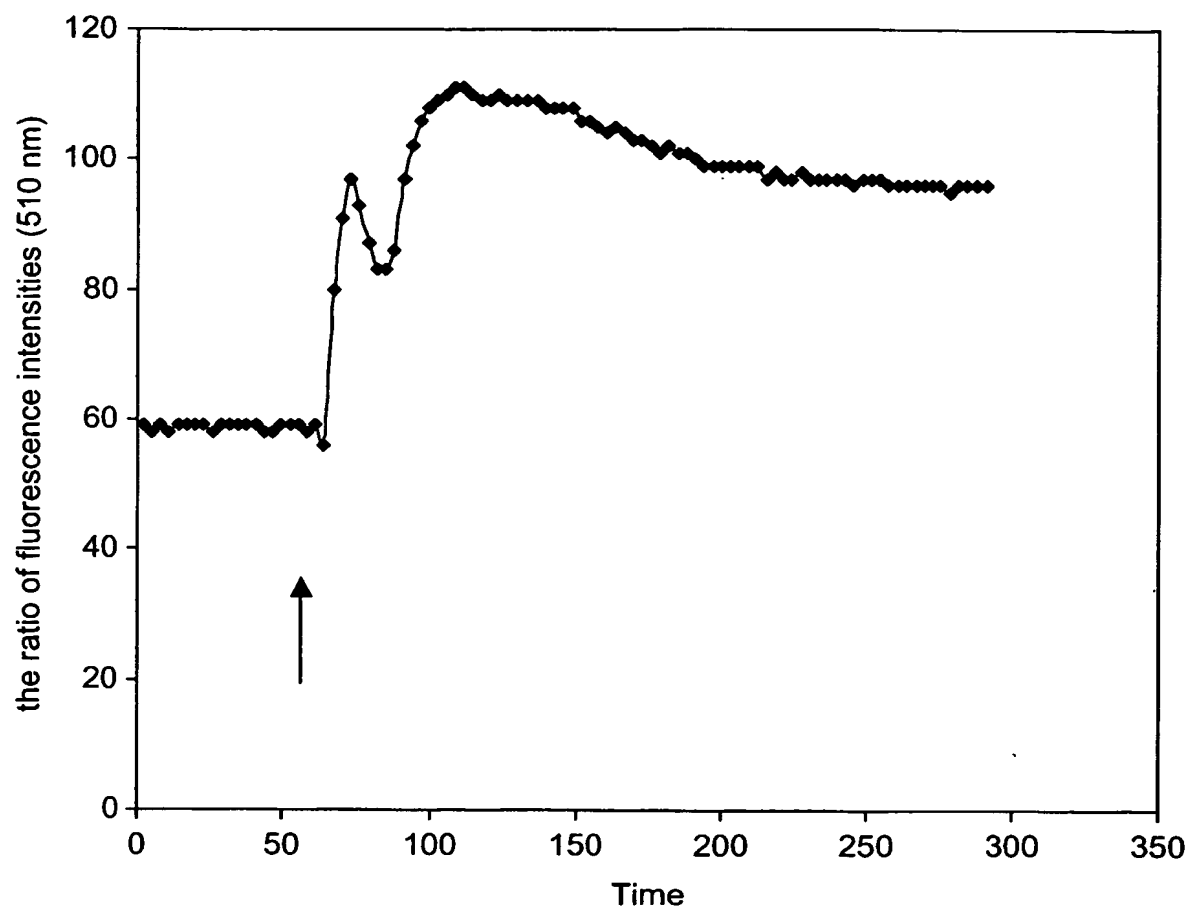


FIG. 6

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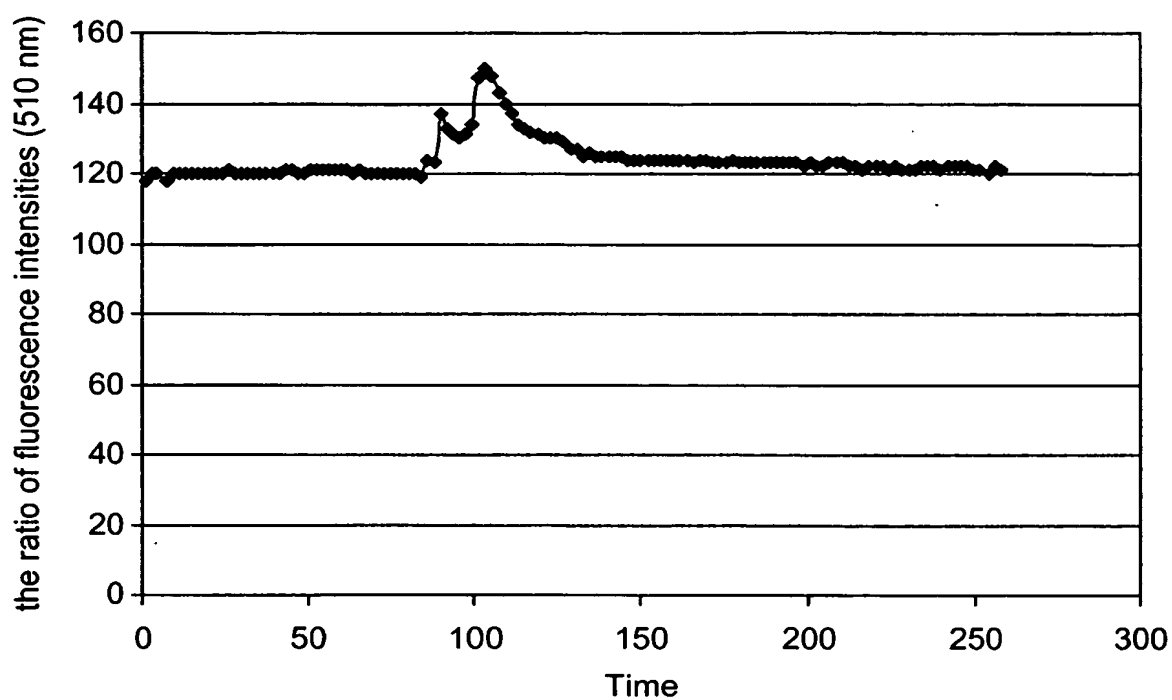


FIG. 7

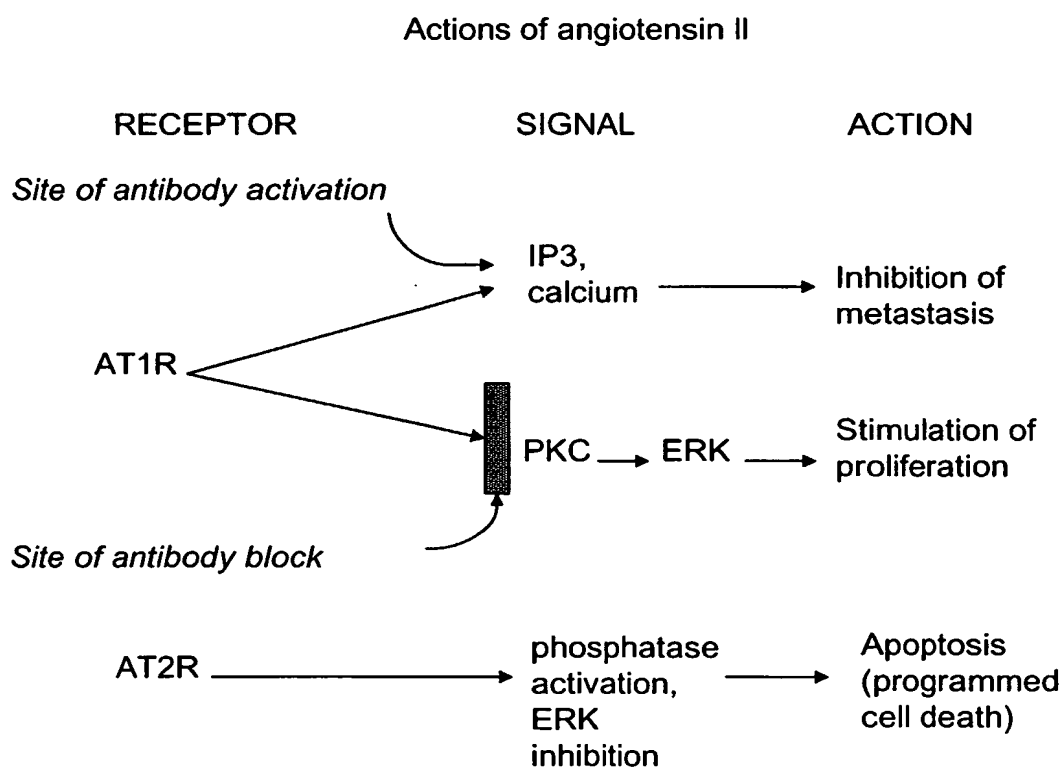


FIG. 8

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| | | | | |
|------------|------|---------|-------------------|-------------------------------|
| Human | AT1 | MILNSST | <u>EDGIKRIQDD</u> | CPKAGRHNHYIFVMIPTLYSIIFVVGIFG |
| Chimp | AT1 | ----- | ----- | ----- |
| Murine | AT1b | ----- | ----- | ----- |
| Bovine | AT1 | ----- | ----- | -----I----- |
| Canine | AT1 | ----- | ----- | ----- |
| Ovine | AT1 | ----- | ----- | -----I-----L-- |
| Rabbit | AT1 | -M----- | ----- | ----- |
| Rat | AT1b | ----- | ----- | -----M----- |
| Guinea pig | AT1 | ----- | Q----- | ---X---S----- |
| Rat | AT1a | -X---X | D----- | -----S----- |
| Mouse | AT1a | -X---X | ----- | --XS--S-----M----- |
| Gerbil | AT1 | -X---X | D----- | -----S----- |

- denotes identical residue

X denotes missing residue

FIG. 9

| | | |
|---|--------------|-------------------------------|
| Applicant's or agent's file reference number | P33791WO/NCB | International application No. |
|---|--------------|-------------------------------|

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

| | |
|---|-------------------------------------|
| A. The indications made below relate to the microorganism referred to in the description on page <u>2</u> , line <u>1 to 9, and on page 10, lines 25 to 29</u> | |
| B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/> | |
| Name of depositary institution European Collection of Animal Cell Cultures (ECACC) | |
| Address of depositary institution (including postal code and country) Vaccine Research and Production Laboratory Public Health Laboratory Service Centre for Applied Microbiology and Research Porton Down, Salisbury Wiltshire SP4 0JG, UK | |
| Date of deposit 22 July 1993 | Accession Number 93072117 |
| C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/> | |
| The deposit was made by Dr Gavin P Vinson and Dr Stewart Barker, Department of Biochemistry, Queen Mary & Westfield College, Mile End Road, London E1 4NS. The depositor has authorised the applicant to refer to the deposited material in the application and has given his unreserved and irrevocable consent to the deposited material being made available to the public in accordance with Rule 28(1)(d) of the European Patent Convention. | |
| D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States) | |
| In respect of all designated States to which such action is possible and to the extent that it is legally permissible under the law of the designated State, it is requested that a sample of the deposited biological material be made available only by the issue thereof to an independent expert, in accordance with the relevant patent legislation, e.g. EPC Rule 28(4), UK Patent Rules 1995, Schedule 2, Paragraph 3, Australian Regulation 3.25(3) and generally similar provisions mutatis mutandis for any other designated State. | |
| E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable) | |
| The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit") | |

| | |
|---|--|
| For receiving Office use only <input type="checkbox"/> This sheet was received with the international application Authorized officer | For International Bureau use only <input type="checkbox"/> This sheet was received by the International Bureau on: Authorized officer |
|---|--|

APPENDIX 3

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BUDAPEST TREATY ON THE INTERNATIONAL
RECOGNITION OF THE DEPOSIT OF MICROORGANISMS
FOR THE PURPOSES OF PATENT PROCEDURE

INTERNATIONAL FORM

TO
DR GAVIN P VINSON & DR STEWART
BARKER
QUEEN MARY & WESTFIELD COLLEGE
DEPARTMENT OF BIOCHEMISTRY
MILE END ROAD
LONDON
E1 4NS
UK

NAME AND ADDRESS
OF DEPOSITOR

| | |
|--|--|
| I. IDENTIFICATION OF THE MICROORGANISM | |
| Identification reference given by the DEPOSITOR: 6313/G2 | Accession number given by the INTERNATIONAL DEPOSITORY AUTHORITY: 93072117 |
| II. SCIENTIFIC DESCRIPTION AND/OR PROPOSED TAXONOMIC DESIGNATION | |
| The microorganism identified under I above was accompanied by: | |
| <input checked="" type="checkbox"/> A scientific description <input type="checkbox"/> A proposed taxonomic designation (Mark with a cross where applicable) | |
| III. RECEIPT AND ACCEPTANCE | |
| This International Depository Authority accepts the microorganism identified under I above, which was received by it on 21 July 1993 (date of the original deposit) ¹ | |
| IV. RECEIPT OF REQUEST FOR CONVERSION | |
| The microorganism identified under I above was received by this International Depository Authority on (date of the original deposit) and A request to convert the original deposit to a deposit under the Budapest Treaty was received by it on (date of receipt of request for conversion) | |
| IV. INTERNATIONAL DEPOSITORY AUTHORITY | |
| Name: Dr D H Lewis Address: ECACC CAMR Porton Down Salisbury SP4 0JG | Signature(s) of person(s) having the power to represent the International Depository Authority or of authorized official(s): Date: 15/12/83 |

¹ Where Rule 6.4(d) applies, such date is the date on which the status of international depository
authority was acquired

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